

Census of the Bacterial Community of the Gypsy Moth Larval Midgut by Using Culturing and Culture-Independent Methods

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Little is known about bacteria associated with Lepidoptera, the large group of mostly phytophagous insects comprising the moths and butterflies. We inventoried the larval midgut bacteria of a polyphagous folivore, the gypsy moth (*Lymantria dispar* L.), whose gut is highly alkaline, by using traditional culturing and culture-independent methods. We also examined the effects of diet on microbial composition. Analysis of individual third-instar larvae revealed a high degree of similarity of microbial composition among insects fed on the same diet. DNA sequence analysis indicated that most of the PCR-amplified 16S rRNA genes belong to the γ -Proteobacteria and low G+C gram-positive divisions and that the cultured members represented more than half of the phylotypes identified. Less frequently detected taxa included members of the α -Proteobacterium, Actinobacterium, and Cytophaga/Flexibacter/Bacteroides divisions. The 16S rRNA gene sequences from 7 of the 15 cultured organisms and 8 of the 9 sequences identified by PCR amplification diverged from previously reported bacterial sequences. The microbial composition of midguts differed substantially among larvae feeding on a sterilized artificial diet, aspen, larch, white oak, or willow. 16S rRNA analysis of cultured isolates indicated that an *Enterococcus* species and culture-independent analysis indicated that an *Entbacter* sp. were both present in all larvae, regardless of the feeding substrate; the sequences of these two phylotypes varied less than 1% among individual insects. These results provide the first comprehensive description of the microbial diversity of a lepidopteran midgut and demonstrate that the plant species in the diet influences the composition of the gut bacterial community.

Microorganisms play important and often essential roles in the growth and development of many insect species. Endosymbionts contribute to insect reproduction, digestion, nutrition, and pheromone production (7, 13, 14, 20, 50, 53, 55). Symbiotic relationships between insects and their gut bacteria have been studied extensively in several systems, particularly in termites and aphids, which feed on wood and plant phloem content, respectively (8, 17, 31). However, relatively little is known about microbial associates in other insect groups, particularly those that feed on foliage.

Describing relationships between insects and their associated bacteria has proved challenging because many of the microbial associates are not readily culturable. The availability of culture-independent tools, such as PCR, provides an opportunity to detect and classify microorganisms that cannot be cultured by existing methods (74). PCR has revealed the phylogeny of endosymbionts and associates of a number of insect species (1, 4, 15, 19, 33, 55, 57, 62, 64).

Little is known about the bacteria associated with Lepidoptera, a primarily phytophagous group that is one of the largest insect orders, containing over 150,000 species (65). A few studies indicate that lepidopterans harbor midgut bacteria (14, 49). These studies suggest the possibility that microorganisms provide essential nutrients or assist in important biochemical functions. However, an understanding of the types of microbes present in the midgut and of their roles in insect development

and function is lacking. Several attributes of Lepidoptera larvae make their bacteria of particular interest. In particular, the high alkalinity (typically pH 8 to 10) of most lepidopteran midguts and the diverse chemistry of the midgut generated by the unusually broad feeding range of some species make this a particularly challenging environment for microorganisms (5, 22, 29, 44). Furthermore, the Lepidoptera include some of the most damaging agricultural and forest pests worldwide, including species that overcome existing control measures, such as synthetic insecticides, microbial insecticides, genetic host plant resistance, and genetically modified plants (60, 70). Knowledge of the gut bacteria of the Lepidoptera and the roles they may play in larval biology could lead to new targets for pest management.

As an initial step toward understanding relationships between Lepidoptera and their midgut bacteria, we surveyed the midgut of the gypsy moth, *Lymantria dispar* (L.), which is an invasive species of Eurasian origin and the most damaging defoliator of deciduous trees in North America (51). The gypsy moth alters natural ecosystems, causes economic losses to the forest industry and homeowners, elicits allergic reactions in humans, and is highly polyphagous with a reported host range of 300 species (41). We examined the effects of diet and insect source on the microbial composition of third-instar midguts.

MATERIALS AND METHODS

Sources and treatments of gypsy moth larvae. (i) **Larvae from diverse populations feeding on a common substrate.** Gypsy moth egg masses were obtained from the culture New Jersey Standard Strain (NJSS) at the U.S. Department of Agriculture (USDA)-Animal and Plant Inspection Service (APHIS) laboratory at the OTIS Air National Guard Base, Cape Cod, Mass. (MA colony) and the

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Beneficial Insects Introduction and Research Laboratory, U.S. Department of Agriculture-Agricultural Research Services, Newark, Del. (DE colony). In addition, egg masses were collected from field sites in Wisconsin (WI field) and Michigan (MI field). Wisconsin egg masses were collected on 24 March 1999 from five trees (four oak trees [*Quercus* sp. L.] and one maple tree [*Acer* sp. L.]) on a rural property in York Township, Dane County (R11E, T9N). The Michigan egg masses were collected 23 March 1999 from a newly established population in a mixed oak and aspen (*Quercus* sp. and *Populus* sp.) stand in Lincoln Township, Clare County (R5W, T18N). Field-collected egg masses were kept in plastic bags at 4°C from the time of collection until they were used in assays. Assays were timed to coincide with Wisconsin field larval emergence (around 5 May 1999).

Gypsy moth larvae were reared as described in Broderick et al. (10). Briefly, egg masses were surface sterilized with a solution of Tween 80 (polyoxyethylene sorbitan monooleate), bleach, and distilled water. Larvae were reared in 17-cm-diameter petri dishes on a sterile artificial diet (USDA Hamden Formula) under a 16 h:8 h (light:dark) photoperiod at 25°C. Colony (MA and DE) larvae were reared in a quarantine facility at the University of Wisconsin—Madison Department of Entomology. To reduce the potential for introduction of pathogens into the quarantine facility, field egg masses and larvae (WI and MI) were reared in a separate campus quarantine facility at the University of Wisconsin—Madison Biotron under the same conditions as above. Larvae were provided with artificial diet after emergence and provided a new diet every 48 h.

(ii) Larvae from a common source feeding on multiple tree species. Gypsy moth egg masses from culture NJSS at the USDA-APHIS laboratory (MA colony) were reared according to methods described previously (10, 11, 18).

Two-year-old white oak (*Q. alba* L.), larch [*Larix laricina* (Du Roi) K. Koch], and quaking aspen (*P. tremuloides* Michaux) trees were obtained from the Wisconsin Department of Natural Resources Nursery, Hayward. Two-year-old scrub willow (*Salix fragilis* L.) trees were obtained from the Iowa State Nursery, Ames. The trees were chilled at 4°C for 20 days after receipt to ensure good bud development. Following the chilling period, the trees were planted in 12-liter pots in Sunshine Mix LCI (Sungro Horticulture, Bellevue, Wash.) and flood irrigated. The trees were grown in a greenhouse at 25°C under a 16 h:8 h (light:dark) photoperiod. Trees were watered every 5 days until bud development; following bud development the trees were watered every 2 to 3 days.

Larvae were provided with a sterilized artificial diet for 24 h after emergence and then were divided into 40 groups of 200 larvae each. These 40 groups were distributed among treatments such that 10 groups of larvae fed on leaves of one of the four tree species in petri dishes lined with a piece of moistened sterilized filter paper. For each tree species 10 trees were sampled, with one group of 200 larvae feeding on leaves collected from each tree (a total of 2,000 larvae for each tree species). Whole leaves were provided, with the petioles inserted in a microcentrifuge tube containing distilled water to prevent desiccation. Foliage was replaced every 48 h until the larvae were used for experiments.

Culture-independent methods. (i) Midgut separation and DNA extraction. Third-instar larvae were surface sterilized for 5 s in 95% ethanol prior to dissection. Dissecting scissors were used to cut laterally behind the head capsule, and the gut was removed from the cuticle with larval forceps. The crop and midgut were collected and placed in a 1.5-ml microcentrifuge tube for processing. Samples were placed in a -80°C freezer.

Gypsy moth midguts were analyzed individually in all experiments. Total microbial DNA was extracted from individual gypsy moth crops and midguts by using a protocol modified from the method described by Ausubel et al. (3). Microcentrifuge tubes, each containing a single gut, were thawed, and 600 µl of Tris-EDTA (TE) (10 mM Tris-HCl [pH 8.0], 1 mM EDTA) was added to each tube. The contents of the tube were then sonicated (50 to 60 Hz, 117 V, 1.0 A; Branson Ultrasonics, Danbury, Conn.) for 30 s to separate bacterial cells from the gut wall, and 537 µl of TE was removed and placed in a new 1.5-ml microcentrifuge tube. The sample was sonicated under the same conditions for 45 s to break open bacterial cells and was mixed thoroughly with 60 µl of 10% sodium dodecyl sulfate and 3 µl of 50 mg of proteinase K/ml and was incubated for 1.5 h at 37°C. Each tube was mixed with 100 µl of 5 M NaCl prior to the addition of 80 µl of 10% cetyltrimethylammonium bromide-5 M NaCl. The sample was mixed thoroughly and incubated at 65°C for 30 min. DNA was extracted with equal volumes of chloroform-isoamyl alcohol (CIA) (24:1 [vol/vol]) and phenol CIA (25:24:1 [vol/vol/vol]). DNA was precipitated with isopropanol and recovered by centrifugation. Pellets were resuspended in 100 µl of TE buffer. DNA concentration was determined by absorbance ratio at 260/280 nm, and the DNA suspension was stored at -20°C until it was used for PCR and further analysis.

(ii) PCR amplification. Bacterial 16S rRNA was amplified by PCR from total DNA by using primers 27F and 1492R (37). For terminal-restriction fragment length polymorphism (T-RFLP) analysis, the 5' primer (27F) was

labeled with the dye 6-carboxyfluorescein (6-FAM) (16, 42). All oligonucleotide primers were synthesized by the Biotechnology Center at the University of Wisconsin—Madison. PCR contents for a 50-µl volume were 25 ng of template DNA, 0.5 µl of 20 µM forward (27F) and reverse (1492R) primers, 0.8 µl of 10 mM deoxynucleoside triphosphates (Gibco BRL, Grand Island, N.Y.), 5 µl of 10× *Taq* polymerase buffer, and 0.5 µl of *Taq* DNA polymerase (2.5 U; Promega, Madison, Wis.). The thermal cycling conditions were an initial 3-min denaturation step at 94°C, 30 cycles at 94°C for 30 s, 55°C for 90 s, and 72°C for 2.5 min, with a final 5-min extension at 72°C. A hot-start protocol (34) was used with *Taq* polymerase added to each reaction after 1 min of the 3-min initial 94°C denaturation step. All reactions were carried out in 0.2-ml reaction tubes in a Robocycler 96 (Stratagene, La Jolla, Calif.). Each PCR was examined by electrophoresis in a 0.8% agarose gel, and gels were visualized by staining with ethidium bromide.

(iii) T-RFLP analysis. Samples of PCR amplification products from 10 individual larvae exposed to each diet type and from each population source were digested (37°C for 3 h) with the restriction endonucleases *AluI* and *HhaI* (Promega). Digestion reactions (10 µl total volume) contained 5 µl of PCR product, 0.3 µl of enzyme, and 0.3 µl of the appropriate incubation buffer according to the synthesis protocol. Samples (5 µl) of the digestion reactions were mixed with 19 µl of deionized formamide and 1 µl of an internal standard (CST ROX 200-2000; BioVentures, Inc., Murfreesboro, Tenn.), denatured for 5 min at 95°C, and kept on ice until they were loaded into an autosampler. The samples were analyzed by capillary electrophoresis in an ABI DNA sequencer (model 310; Perkin-Elmer, Foster City, Calif.) (39). Data from individual samples were analyzed with GeneScan 3.1 (Perkin-Elmer) and were compared to other samples by using the Ribosomal Database Project II (RDP II) T-RFLP Online Analysis software (46). To assess the similarity of microbial composition among midguts, T-RFLP profiles from replicate samples were aligned and analyzed by pairwise comparison (40). Similarity values, S_{ab} , were determined by using the equation $2N_{ab} / (N_a + N_b)$, where N_{ab} is the number of peaks in common between samples and N_a and N_b are the number of total peaks in each sample.

(iv) Cloning 16S rRNA genes. Total DNA from pools of 10 larvae from all population and diet treatment samples was PCR-amplified separately by using the bacterial primers 27F and 1492R. Clone libraries were constructed by ligating PCR product into pGEM-T Vector system (Promega) according to the manufacturer's instructions and by transforming competent *Escherichia coli* cells (DH5α) by electroporation (61). Plasmid DNA was isolated from transformants by using QIAprep Spin Plasmid Miniprep kits (QIAGEN, Valencia, Calif.). All clones in libraries of approximately 250 clones from each diet type were sequenced.

(v) 16S rRNA sequencing and data analysis. Sequencing reactions were performed by using the BigDye reaction mix (Perkin-Elmer Corp.) according to the manufacturer's instructions. Purified plasmid DNA was initially sequenced by using the plasmid primers T7 and SP6, which flank the insert DNA in pGEM-T, and additionally sequenced with 704F and 787R to obtain full 1,500-bp sequences (37). DNA from cultured organisms was purified by using the PCR product Pre-sequencing Kit (USB, Cleveland, Ohio) and was sequenced by using either 27F, 1492R, 704F, or 787R (37). Two representatives of each colony morphology and a total of 400 clones (approximately 45 to 115 per diet treatment) were analyzed. All 16S rRNA sequences (1,450 to 1,490 bp) were compiled by using the SeqMan program from the DNASTar software package (DNASTAR, Inc., Madison, Wis.) and were compared to available databases by use of BLAST to determine approximate phylogenetic affiliations (2). Individual sequences of isolates and clones that were ≥98% identical to each other were considered the same phylotype and were combined for analysis. All sequences were tested for possible chimeric structures by using RDP Check_Chimera (39).

(vi) Culturing and enumeration of midgut bacteria. Twenty-third-instar larvae from each of the five diet treatments were dissected for culturing studies following a 24-h starvation period to allow gut clearing. Midguts were each placed in 1 ml of phosphate-buffered solution (PBS) in a 1.5-ml microcentrifuge tube. Ten larvae feeding on foliage from two trees of each species were used. Individual guts in PBS were sonicated for 45 s and then were diluted plated on 1/10 strength tryptic soy agar (TSA) at neutral pH and were incubated at 28°C. Colonies were classified based on morphological parameters of shape, color, margins, elevation, and texture. Colonies of each designated morphology were counted after 24, 48, and 72 h. Isolates of two representatives of each colony morphology were purified for individual colonies, and cellular morphology of each isolate was examined with 1,000× magnification. DNA was extracted with the EasyDNA kit (Invitrogen, Carlsbad, Ohio) from 14-h-old cultures of each isolate in 1/2-strength tryptic soy broth. DNA was quantified, amplified, and sequenced as described above.

Ten individual third-instar midguts from larvae feeding on artificial diet were

TABLE 1. Bacterial phylotypes identified by culturing and culture-independent analysis of third-instar gypsy moth midguts based on 16S rRNA gene sequence analysis

Sequence identity and identification method	Bacterial division	Genus ($\geq 95\%$ identity)	Database matches ($\geq 98\%$ identity)	Detection in gypsy moth fed on ^a :				
				Artificial diet	Larch	Willow	Aspen	White oak
Identified by culturing and culture-independent analysis								
NAB1	γ -Proteobacteria	<i>Pseudomonas</i>	<i>P. putida</i>	1.8×10^5	1.1×10^7		3.6×10^5	3.5×10^5
NAB2	γ -Proteobacteria	<i>Pseudomonas</i>			2.2×10^6			
NAB3	γ -Proteobacteria	<i>Enterobacter</i>	Uncultured soil bacterium F42326.1	4.8×10^6	7.0×10^7	8.9×10^7		
NAB4	γ -Proteobacteria	<i>Pantoea</i>	<i>P. agglomerans</i>	2.5×10^5	3.7×10^4	1.5×10^7		
NAB5	γ -Proteobacteria	<i>Serratia</i>	<i>S. marcescens</i>		4.6×10^4		7.5×10^8	4.0×10^4
NAB6	low G+C gram positive	<i>Bacillus</i>			4.0×10^8			
NAB7	low G+C gram positive	<i>Staphylococcus</i>	<i>S. lentus</i>	8.1×10^4		9.1×10^6		4.5×10^6
NAB8	low G+C gram positive	<i>Staphylococcus</i>	<i>S. cohnii</i>	2.9×10^6	4.7×10^6	1.3×10^7		
NAB9	low G+C gram positive	<i>Staphylococcus</i>	<i>S. xylosus</i>	1.3×10^6	1.07×10^5			
NAB10	low G+C gram positive	<i>Paenibacillus</i>			2.6×10^6			
NAB11	low G+C gram positive	<i>Enterococcus</i>	<i>E. faecalis</i>	1.5×10^8	1.9×10^6	3.6×10^8	1.4×10^4	2.1×10^7
NAB12	low G+C gram positive	<i>Enterococcus</i>				4.8×10^7	4.1×10^6	2.4×10^5
NAB13	Actinobacterium group	<i>Rhodococcus</i>				1.2×10^7		
NAB14	Actinobacterium group	<i>Microbacterium</i>			5.2×10^8		2.2×10^6	
NAB15	CFB ^b group						7.4×10^4	
Identified by culture-independent analysis only								
NAB16	α -Proteobacteria	<i>Agrobacterium</i>		+		+	+	
NAB17	γ -Proteobacteria	<i>Enterobacter</i>		+	+	+	+	+
NAB3	γ -Proteobacteria	<i>Enterobacter</i>			+			
NAB18	γ -Proteobacteria				+		+	
NAB19	γ -Proteobacteria				+			+
NAB20	low G+C gram positive			+	+	+		
NAB21	Actinobacterium group	<i>Micrococcus</i>					+	
NAB22	Actinobacterium group						+	
NAB23	CFB group						+	

^a For cultured cells the numbers represent the average population of phylotype based on colony morphology in 20 individual larvae per treatment (CFU/milliliter of gut material). For noncultured samples, a plus indicates presence in guts of insects in diet treatment.

^b CFB, *Cytophaga/Flavobacteria/Bacteroides*.

dissected, sonicated, and stained with 4,6-diamino-2-phenylindole (DAPI) to determine the total number of bacterial cells per gut (72). For each individual midgut, twenty randomly selected fields were counted by epifluorescence microscopy (Olympus America Inc., Melville, N.Y.).

Nucleotide sequence accession numbers. The compiled 16S rRNA sequences have been deposited in the GenBank database under accession numbers AY39005 to AY39035.

RESULTS

Bacterial diversity in third-instar gypsy moth midguts. Evaluation of the midgut bacteria of gypsy moth larvae using a combination of 16S rRNA analysis of the cultured bacteria and culture-independent PCR amplification of 16S rRNA sequences led to the identification of 23 phylotypes (Table 1). Approximately 65% of these phylotypes are newly reported sequences (<98% identity to any of the bacterial 16S rRNA sequences within GenBank). The majority (70%) of these bacterial sequences fell within the low G+C gram-positive and γ -Proteobacteria divisions. *Actinobacterium*, *Cytophaga/Flexibacter/Bacteroides*, and α -Proteobacteria accounted for 17, 9, and 4% of the sequences, respectively.

Variation in bacterial composition among individual larvae. Comparisons of the T-RFLP profiles of the bacteria in replicate larvae from either the same population source or the same diet treatment showed a high similarity index of between 0.96 and 1.00 (Tables 2 and 3). These kinds of similarities were also reflected in the types of colony morphologies obtained in the culture-dependent analyses. The majority of isolates examined were found in >90% of the larvae examined and all were isolated from >50% of the larvae in a given treatment.

To identify factors associated with variation in microbial composition, we compared PCR-amplified 16S rRNA genes isolated from the midguts of gypsy moth from lab and field populations by using T-RFLP analysis. The T-RFLP profiles of the MA colony and WI field samples and those of the DE colony and MI field all fed artificial diet were very similar (Table 2). Moreover, these pairs differed only slightly (less than 6%) from each other (Table 2), indicating that egg source did not have a significant effect on the bacteria of the gypsy moth midgut. Given these similarities, only the MA colony population was used to examine the variation in bacterial di-

TABLE 2. Pairwise comparisons for similarity of T-RFLPs from the midgut of gypsy moth from lab and field populations feeding on an artificial diet

Population	Coefficient within population	Coefficient between populations		
		MA colony	DE colony	WI field
MA colony	0.996			
DE colony	0.997	0.952		
WI field	0.996	0.994	0.958	
MI field	0.998	0.955	0.992	0.953

versity due to diet. Diet greatly affected the T-RFLP profiles (Fig. 1), as indicated by the lower similarity values of T-RFLP profiles from different diets (Table 3).

Comparison of bacterial diversity revealed by culturing and culture-independent analysis. Culturing yielded 15 distinguishable types of bacteria across all diet treatments from gypsy moth midguts based on cellular and colony morphological traits; 93% of these isolates could be assigned to a genus, while only 47% could be assigned to a species by using 16S rRNA analysis (Table 1). DAPI staining and microscopic enumeration indicated that there were 4×10^8 viable bacterial cells/gut. The phylotypes indicated by culture-independent methods exhibited greater divergence and diversity than phylotypes recovered by culturing. In addition to the 15 phylotypes recovered by culturing, 9 phylotypes were indicated based on 16S rRNA analysis in gypsy moth midguts across all diet treatments; only four of these phylotypes could be assigned definitively to a genus.

Effect of diet on bacterial composition. Two phylotypes, *Enterococcus faecalis* and an uncultivated *Enterobacter* sp. (NAB17) were found in all larvae, regardless of treatment. In comparing the relative diversity of gut bacteria associated with each diet treatment, we found that 8 bacterial phylotypes were detected in only one of the diet treatments (Table 1). Larvae feeding on larch contained the greatest diversity of bacterial phylotypes (15, of which 11 were cultivated). Larvae feeding on aspen contained the highest proportion of uncultivated bacterial phylotypes of all the diet treatments. Among the 13 phylotypes identified from this group, over 50% of the phylotypes were obtained by using culture-independent methods only. On average, 51% of phylotypes in each diet treatment were previously reported sequences.

Comparison of T-RFLP profiles and 16S rRNA sequences. To assess the ability of T-RFLP analysis to describe the bac-

terial composition of the midgut, 16S rRNA sequences derived from midguts were analyzed *in silico* by using the RDP II T-RFLP Analysis Program (43) and were compared to actual T-RFLPs from the same samples. In general, 16S rRNA sequences showed the predicted peak pattern, and all sequences could be accounted for in the T-RFLP. However, this analysis also demonstrated that multiple fragments were found at the same peak position regardless of the restriction enzyme, especially among similar 16S sequences. These results are consistent with previous indications of the resolution limits of T-RFLP analysis in assessing the full extent of diversity (23). However, these results also indicate that T-RFLP analysis is useful as a first step to assess sample diversity and identify treatments or conditions for further investigation.

DISCUSSION

These results demonstrate that the microbial diversity of the gypsy moth midgut is relatively simple, with 15 phylotypes at its most complex and 7 phylotypes at its simplest. The relative simplicity of this community is especially apparent compared to other gut environments, such as the human intestine and termite hindguts, which contain at least 500 and 50 phylotypes, respectively (54, 76). We expect that the census presented here is fairly complete given that we can account for more than 40% of the viable cells by culturing techniques, and all of the peaks in the T-RFLP analysis are accounted for in the PCR-generated 16S rRNA sequences. However, there may well be minor species whose presence is obscured because they were not cultured by our methods and were not detected by either T-RFLP analysis or 16S rRNA sequence analysis. Diet had a significant impact on midgut microbial diversity, as shown by culturing and sequence analysis of 16S rRNA genes amplified directly from larval midguts. Molecular, culture-independent methods revealed that many of the amplified 16S rRNA genes from gypsy moth midguts consisted of sequences not previously described. Perhaps the most surprising result of this study was the culturing of numerous bacterial species whose 16S rRNA sequences were as yet undescribed in public databases and, in some cases, diverged deeply from known species or genera.

Molecular characterization of cultured isolates and culture-independent methods identified *E. faecalis* in all midguts. While the 16S rRNA sequences from these isolates match most closely to *E. faecalis*, preliminary analysis revealed that these isolates differ physiologically (data not shown) from the clinical isolates of *E. faecalis* reported previously (21, 52). However, other *E. faecalis* isolates from environmental samples consistently diverge from the clinical type strains in certain physiological characteristics, including colony color, carbohydrate metabolism profiles, and growth temperature profiles (26, 47). Further analysis is required to determine the implications of these differences for species definition within the genus *Enterococcus* (63).

The novelty of phylotypes found in the gypsy moth midguts was anticipated, based on the insect's physiology. While the caterpillar lives in a common terrestrial macroenvironment as an external phytophage, the microenvironment in which these midgut bacteria reside is chemically extreme. Most insect mid-

TABLE 3. Pairwise comparisons for similarity of T-RFLPs from the midgut of gypsy moth fed various diets

Diet	Coefficient within group	Coefficient between diet groups			
		Artificial diet	Larch	Aspen	Willow
Artificial diet	0.993				
Larch	0.962	0.382			
Aspen	0.975	0.139	0.029		
Willow	0.968	0.077	0.053	0.083	
White Oak	0.989	0.045	0.028	0.038	0.091

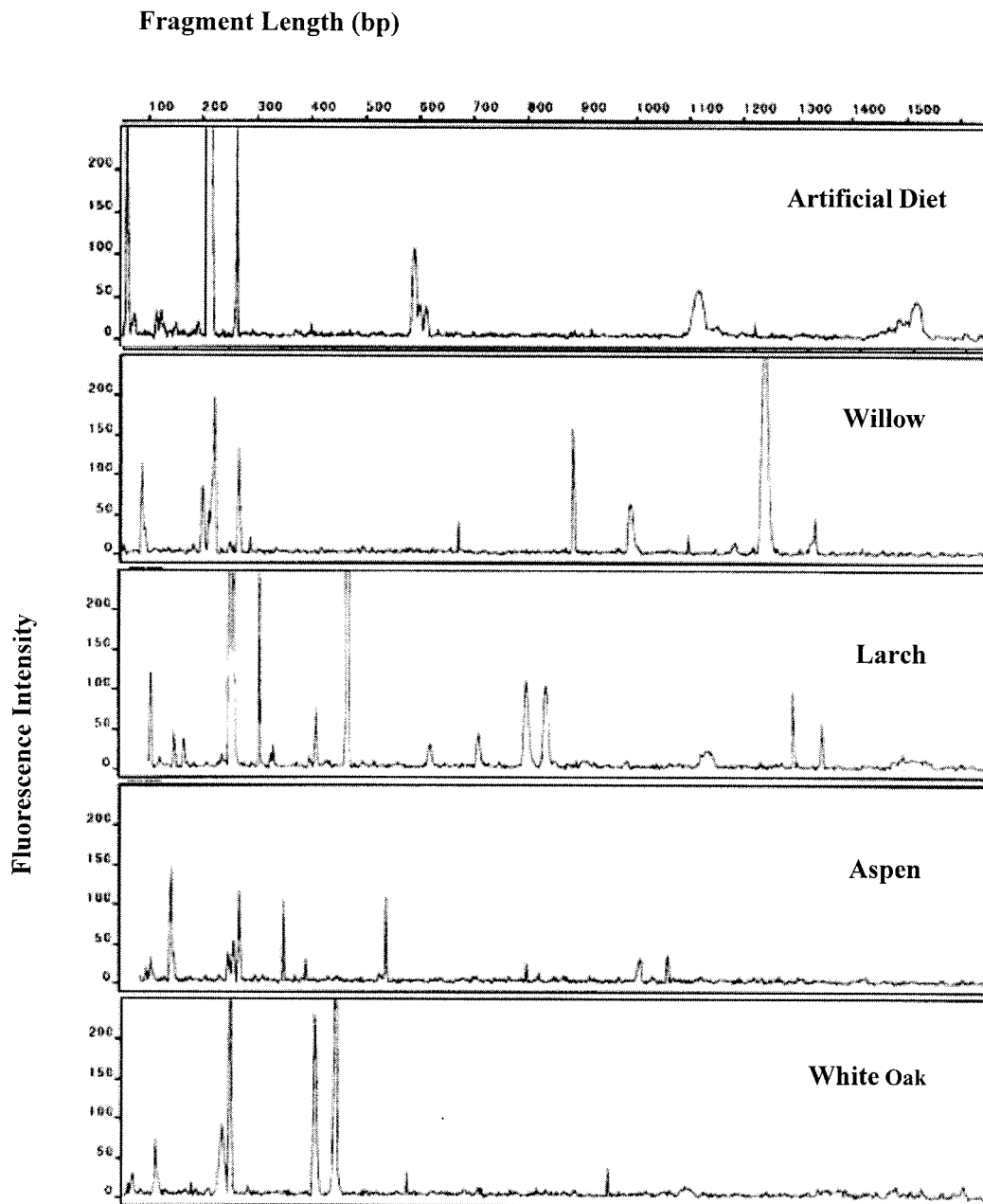


FIG. 1. T-RFLP profiles of third-instar midguts of gypsy moth feeding on diverse diets. Each profile is representative of 10 individual larvae.

guts are generally neutral to acidic (pH 4 to 7), while lepidopteran midguts are typically pH 8 to 10. However, values as high as 12.4 have been reported for gypsy moth (5, 22, 29). In addition, the diverse plants on which this insect feeds contain an array of allelochemicals, such as phenolics, tannins, and terpenoids, many of which are toxic to microorganisms (28). Therefore, this extreme environment may have selected for unusual phenotypes, resulting in unusual phylotypes.

The microbial midgut inhabitants seem likely to benefit the insect host, based on observations of other insect systems (7, 9, 14, 31). Examples of such benefits may include modifying mid-

gut pH, detoxifying plant allelochemicals, and maintaining the midgut microbial community structure. Midgut pH is important from the perspective of both the microbes and the insect. A predominant member of the midgut microbial community, *E. faecalis*, is commonly found at higher pH (8 to 9) and acidifies its environment through its metabolism (45). This could confer some advantage to the insect host, as some microbial toxins of lepidoptera, such as *Bacillus thuringiensis* toxin, are activated by alkaline conditions (75). Thus, *E. faecalis* might help protect this insect from *B. thuringiensis* toxin by decreasing the midgut pH. Previous work has shown that lar-

vae that are more susceptible to *B. thuringiensis* toxin have smaller populations of *E. faecalis* in their midguts (11). Bacteria may also contribute to their own survival by managing midgut pH to enhance their own growth or exclude competing microorganisms. While numerous studies have investigated microbial diversity in neutral and acidic environments (6, 24, 27, 32, 56, 66), few have examined bacteria in highly alkaline environments, particularly in nonaquatic ecosystems (25, 30). Further analysis of these bacteria may identify adaptations that permit them to function in this extreme environment, and whether they have a role in maintaining it.

The broad range of phytochemicals consumed by this polyphagous herbivore may present a challenge for both the gypsy moth and its associated bacteria. Toxic compounds may select for bacteria that can metabolize them, and these bacteria may degrade ingested compounds that are otherwise toxic to the insect (41). For example, the 16S rRNA gene sequence of one cultured isolate from the gypsy moth (NAB13) was most similar to a *Rhodococcus* species, and recent work identified an enzyme produced by *R. erythropolis* DCL14 (73) that degrades monoterpenes, a widespread class of phytochemicals that can be toxic to many insect larvae and adults (38). The gypsy moth shows relatively higher tolerance to monoterpenes than many other insects (58). If the gypsy moth isolate of *Rhodococcus* species also produces this enzyme, it might contribute to the detoxification of monoterpenes. Identifying traits that facilitate bacterial survival in the gypsy moth midgut may lead to novel chemistry and degradation processes that allow bacteria to live in other extreme environments.

Establishment and maintenance of microbial assemblages within the larval midgut may contribute to insect health, perhaps by suppression of pathogens. For example, one cultured isolate from the gypsy moth midgut (NAB5) is highly similar (>99%) to a sequence found in the gut of the sheep mite, *Psoroptes ovis* (48), and both match *Serratia marcescens* to the species level (>98%). Similar sequences have been found in most insects and other arthropods (12, 35, 67). *Serratia* species are generally considered pathogens and can cause disease in insect production facilities (35, 68). The frequency at which this bacterium occurs across diverse arthropods suggests an important relationship with members of this phylum. *Serratia* might be an opportunistic pathogen living in the insect midgut at subclinical levels, causing disease only when the community structure is altered. The bacteria in the community may also contribute to each others' survival by providing nutrients or processing wastes.

Despite the relatively simple composition of the gypsy moth midgut community, there are challenges to elucidating the roles of various bacteria in gypsy moth biology and in maintenance of species composition within the midgut. In particular, there are several possibilities concerning the establishment of the bacterial midgut composition. For example, it is not known whether midgut bacteria are established through ingestion of plant material and whether different host plant chemistries select for specific bacterial genotypes. The plants selected for this study represent a diversity of secondary plant metabolites that the gypsy moth commonly encounters in the field and may contribute to the observed differences in the midgut bacterial composition. In addition, individual gypsy moth larvae routinely switch host plants in the field with only minor direct

effects on the larvae (69, 70). However, host plant switching can alter the performance of gypsy moth parasitoids and could likewise affect the composition of the bacterial community (36). The relative stability of the gut bacterial community is also unknown. Further analysis of this system, including identification of the source of microbial inoculum in insects fed on sterilized artificial diet, use of methods, such as metagenomics (59), that link function to phylogeny, and improved culturing techniques will advance our understanding of the role of microbes in the gypsy moth midgut.

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